

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 6 :

H01M 10/36**A1**

(11) International Publication Number:

WO 98/18173

(43) International Publication Date:

30 April 1998 (30.04.98)(21) International Application Number: **PCT/US97/18240**(22) International Filing Date: **20 October 1997 (20.10.97)**

(30) Priority Data:

08/735,879**23 October 1996 (23.10.96)****US**(71) Applicant (for all designated States except US): **VALENCE TECHNOLOGY, INC. [US/US]; 301 Conestoga Way, Henderson, NV 89015 (US).**

(72) Inventors; and

(75) Inventors/Applicants (for US only): **BRODD, Ralph [US/US]; 2161 Fountain Springs Drive, Henderson, NV 89014 (US). CHALONER-GILL, Benjamin [US/US]; 520 Mansion Court #306, Santa Clara, CA 95054 (US). GOLOVIN, Milton, Neal [US/US]; 3416 Stillbrook Way, Marietta, GA 30062 (US). ISAACSON, Mark [US/US]; 513 Hickory Place, Santa Clara, CA 95051 (US). LUNDQUIST, Joseph [US/US]; 198 Crystal Lane, Aurora, OH 44202 (US).**(74) Agents: **KREBS, Robert, E. et al.; Burns, Doane, Swecker & Mathis, L.L.P., P.O. Box 1404, Alexandria, VA 22313-1404 (US).**(81) Designated States: **AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).****Published***With international search report.**Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.*(54) Title: **SOLID ELECTROLYTES CONTAINING TOUGHENING AGENTS AND ELECTROLYTIC CELLS PRODUCED THEREFROM**

(57) Abstract

A solid electrolyte containing a polymeric matrix, a salt, a solvent, and a toughening agent, is provided. The toughening agent may include alumina, silica, zeolite, and metal oxides (e.g., calcium oxide and magnesium oxide), and mixtures thereof. The toughening agent acts as a drying agent to remove excess solvent in the electrolyte. The solid electrolytes have improved mechanical strength and adherence to the anode and cathode.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece			TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	NZ	New Zealand		
CM	Cameroon			PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakhstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	SG	Singapore		

SOLID ELECTROLYTES CONTAINING TOUGHENING AGENTS AND ELECTROLYTIC CELLS PRODUCED THEREFROM

BACKGROUND OF THE INVENTION

Field of the Invention

This invention is directed to solid electrolytes containing additives including a toughening agent and/or a basic sink material and, in particular, to solid electrolytes containing a polymeric matrix, a salt, a solvent, and said additives. The toughening agent imparts mechanical strength to the solid electrolyte and the basic sink removes or traps acids (e.g., HF) in the solid electrolyte.

State of the Art

Electrolytic cells containing an anode, a cathode and a solid, solvent-containing electrolyte incorporating a salt are known in the art and are usually referred to as "solid batteries". See, for instance, U.S. Patent Nos. 5,229,225, 5,238,758, 5,358,801, and 5,366,928. These cells offer a number of advantages over electrolytic cells containing a liquid electrolyte (i.e., "liquid batteries") including improved safety features.

Solid batteries employ a solid electrolyte interposed between a cathode and an anode. The solid electrolyte contains either an inorganic or an organic matrix and a suitable salt, such as an inorganic ion salt, as a separate component. Electrolytic cells containing a solid electrolyte having a polymeric matrix suffer from low ion conductivity and, accordingly, in order to maximize the conductivity of these materials, the matrix is generally constructed as a very thin film, i.e., in the range of about 25 to about 250 μm . Minimizing the thickness of the film reduces the total amount of internal resistance within the electrolyte but also decreases the solid electrolyte's structural integrity. In

addition, good adherence of the anode and cathode to the solid electrolyte is necessary for optimum operation of electrochemical cells made therefrom.

Another problem encountered in electrolytic cells is the presence of impurities such as acids (e.g., HF) in the solid and liquid electrolytes. HF is derived from certain lithium salts (e.g., LiPF_6) that are employed. For example, LiPF_6 reacts with water to form HF, LiF (insoluble) and other by-products, thereby reducing the amount of salt available. The acids adversely effect electrochemical performance.

In view of the above, the art is in need of solid electrolytes having superior mechanical attributes, including toughness, hardness, and resiliency. In addition, the solid electrolyte should also adhere to the anode and cathode layers of the electrolytic cell to minimize internal resistance and increase electrochemical performance. Furthermore, there is a need to reduce or eliminate acidic impurities in the solid electrolyte.

SUMMARY OF THE INVENTION

The present invention is based, in part, to the discovery that adding a toughening agent to an electrolyte composition yields a stronger solid electrolyte that adheres well to the anode and cathode. Suitable toughening agents comprise alumina, silica, zeolites, metal oxides and mixtures thereof. The invention is also based in part on the discovery that alumina (Al_2O_3) acts as a base sink to remove or trap acidic impurities, especially HF.

In one aspect, the present invention is directed to an electrolytic cell which comprises: an anode; a cathode and a solid, solvent-containing electrolyte which comprises a polymeric matrix, a salt, a solvent, and a toughening agent, wherein the electrolyte is interposed between the anode and cathode.

In another aspect, the present invention is directed to a process for preparing the electrolytic cells which demonstrate improved electrochemical performance.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

5 This invention is directed to a solid, solvent-containing electrolyte comprising a toughening agent. One aspect of the invention is that the presence of the toughening agent enhances the mechanical strength of the solid electrolyte. Another aspect is that the solid electrolyte layer also demonstrates good adherence to the cathode and/or anode layers. It is believed that
10 improvement in the structural integrity of the solid electrolyte is due, in part, to the ability of the toughening agent to also function as an absorbent (i.e., drying agent) to remove water, excess solvents and impurities. In particular, alumina, a toughening agent, also functions as a basic sink to remove a significant part of the acids from the solid electrolyte which improves cell performance and
15 cycle life.

However, prior to describing this invention in further detail, the following terms are defined as follows:

The term "solid polymeric matrix" refers to an electrolyte compatible material formed by polymerizing organic monomers (or partial polymers thereof) and which, when used in combination with the other components of the electrolyte, renders the electrolyte solid.
20

The term "a solid matrix forming monomer" refers to organic materials which in monomeric form can be polymerized, preferably in the presence of an inorganic ion salt, and a solvent to form solid matrices which are suitable for use as solid elect. in electrolytic cells. Examples of suitable organic solid
25 matrix forming monomers include, for example, propylene oxide,

ethyleneimine, ethylene oxide, epichlorohydrin, acryloyl-derivatized polyalkylene oxides (as disclosed in U.S. Patent No. 4,908,283), urethane acrylate, vinyl sulfonate polyalkylene oxides (as disclosed in U.S. Patent No. 5,262,253, which patent is incorporated herein by reference in its entirety), and
5 the like as well as mixtures thereof.

The term "a partial polymer of a solid matrix forming monomer" refers to solid matrix forming monomers which have been partially polymerized to form reactive oligomers. Partial polymerization may be conducted for the purpose of enhancing the viscosity of the monomer, decreasing the volatility of
10 the monomer, and the like. Partial polymerization is generally permitted so long as the resulting partial polymer can be further polymerized, preferably in the presence of a solvent, such as, a mixture of organic carbonate(s) to form solid polymeric matrices which are suitable for use as solid electrolytes in electrolytic cells.

15 The term "cured" or "cured product" refers to the treatment of the solid matrix forming monomer or partial polymer thereof under polymerization conditions (including cross-linking) so as to form a solid polymeric matrix. Suitable polymerization conditions are well known in the art and include by way of example, heating the monomer, irradiating the monomer with UV light,
20 electron beams, and the like. Examples of suitable cured products suitable for use in this invention are set forth in U.S. Patent Nos. 4,830,939 and 4,990,413 which are incorporated herein by reference in their entirety.

The solid matrix forming monomer or partial polymer can be cured or further cured prior to or after addition of the salt, solvent, and toughening
25 agent, and, optionally, a viscosifier. For example, a composition comprising requisite amounts of the solid matrix forming monomer, salt, organic carbonate solvent, viscosifier and toughening agent can be applied to a substrate and then

cured. Alternatively, the matrix forming monomer can be first cured and then dissolved in a suitable volatile solvent. Requisite amounts of the salt, organic carbonate solvent, viscosifier and toughening agent can then be added. The mixture is then placed on a substrate; removal of the volatile solvent would
5 result in the formation of a solid electrolyte. In either case, the resulting solid electrolyte would be a homogeneous, single phase product which is maintained upon curing, and does not readily separate upon cooling to temperatures below room temperature. Accordingly, the solid electrolyte of this invention does not require a separator as is typical of liquid electrolytes.

10 Alternatively, the solid polymeric matrix can be formed by a casting process which does not require the use of monomers or prepolymers, that is, no curing is required. A preferred method employs a copolymer of polyvinylidenedifluoride and hexafluoropropylene dissolved in acetone or other
15 suitable solvent(s). Upon casting the solution, the solvent is evaporated to form the solid polymeric matrix. The solution may be casted directly onto a current collector. Alternatively, the solution is casted onto a substrate, such as a carrier web, and after the solvent (e.g., acetone) is removed, an electrode film is formed thereon. Preferably, the toughening agent and/or basic sink material is incorporated into the solid polymeric matrix by adding the same to the
20 solution prior to casting. Casting techniques of preparing electrolytic cells are described for in U.S. Patent application serial number 08/559,121 filed on November 17, 1995, which application is incorporated herein.

The term "salt" refers to any salt, for example, an inorganic salt, which is suitable for use in a solid electrolyte. Representative examples of suitable
25 inorganic ion salts are alkali metal salts of less mobile anions of weak bases having a large anionic radius. Examples of such anions are I^- , Br^- , SCN^- , ClO_4^- , BF_4^- , PF_6^- , AsF_6^- , CF_3COO^- , $CF_3SO_3^-$, $N(SO_2CF_3)_2^-$, and the like. Specific examples of suitable inorganic ion salts include $LiClO_4$, $LiSCN$, $LiBF_4$, $LiAsF_6$,

LiCF₃SO₃, LiPF₆, (CF₃SO₂)₂NLi, (CF₃SO₂)₃CLi, NaSCN, and the like. The inorganic ion salt preferably contains at least one cation selected from the group consisting of Li, Na, Cs, Rb, Ag, Cu, Mg and K. When employing salts such as, for example, LiBF₄, LiPF₆, or LiAsF₆, which forms HF in the liquid and solid electrolytes, preferably a base sink material such as alumina is also added to the solid electrolyte to neutralize the HF.

The term "compatible electrolyte solvent" or "electrolytic solvent," or in the context of components of the solid electrolyte, just "solvent," is a low molecular weight organic solvent added to the electrolyte and/or the cathode composition, which may also serve the purpose of solvating the inorganic ion salt. The solvent is any compatible, relatively non-volatile, aprotic, relatively polar, solvent. Preferably, these materials have boiling points greater than about 85°C to simplify manufacture and increase the shelf life of the electrolyte/battery. Typical examples of solvent are mixtures of such materials as propylene carbonate, ethylene carbonate, dimethyl carbonate, diethyl carbonate, gamma-butyrolactone, triglyme, tetraglyme, dimethylsulfoxide, dioxolane, sulfolane, and the like. A particularly preferred solvent is disclosed in U.S. Patent 5,262,253, which is incorporated herein solvent is disclosed in U.S. Patent 5,262,253, which is incorporated herein.

For electrochemical cells where the cathode comprises lithiated cobalt oxides, lithiated manganese oxides, or lithiated nickel oxides, and where the anode comprises carbon, the electrolytic solvent preferably comprises a mixture of ethylene carbonate and dimethyl carbonate. For electrochemical cells where the cathode comprises vanadium oxides, e.g., V₆O₁₃ and the anode is lithium, the electrolytic solvent preferably comprises a mixture of propylene carbonate and triglyme.

The term "organic carbonate" refers to hydrocarbyl carbonate

compounds of no more than about 12 carbon atoms and which do not contain any hydroxyl groups. Preferably, the organic carbonate is a linear aliphatic carbonate and most preferably a cyclic aliphatic carbonate.

Suitable cyclic aliphatic carbonates for use in this invention include

5 1,3-dioxolan-2-one (ethylene carbonate); 4-methyl-1,3-dioxolan-2-one (propylene carbonate); 4,5-dimethyl-1,3-dioxolan-2-one; 4-ethyl-1,3-dioxolan-2-one; 4,4-dimethyl-1,3-dioxolan-2-one; 4-methyl-5-ethyl-1,3-dioxolan-2-one; 4,5-diethyl-1,3-dioxolan-2-one; 4,4-diethyl-1,3-dioxolan-2-one; 1,3-dioxan-2-one; 4,4-dimethyl-1,3-dioxan-2-one; 5,5-dimethyl-1,3-dioxan-2-one; 5-methyl-1,3-dioxan-2-one; 4-methyl-1,3-dioxan-2-one; 5,5-diethyl-1,3-dioxan-2-one; 4,6-dimethyl-1,3-dioxan-2-one; 4,4,6-trimethyl-1,3-dioxan-2-one; and spiro (1,3-oxa-2-cyclohexanone-5',5',1',3'-oxa-2'-cyclohexanone).

10

Several of these cyclic aliphatic carbonates are commercially available

15 such as propylene carbonate and ethylene carbonate. Alternatively, the cyclic aliphatic carbonates can be readily prepared by well known reactions. For example, reaction of phosgene with a suitable alkane- α,β -diol (dihydroxy alkanes having hydroxyl substituents on adjacent carbon atoms) or an alkane- α,γ -diol (dihydroxy alkanes having hydroxyl substituents on carbon atoms in a 1,3 relationship) yields an a cyclic aliphatic carbonate for use within

20 the scope of this invention. See, for instance, U.S. Patent No. 4,115,206, which is incorporated herein by reference in its entirety.

Likewise, the cyclic aliphatic carbonates useful for this invention may be prepared by transesterification of a suitable alkane- α,β -diol or an alkane- α,γ -diol with, e.g., diethyl carbonate under transesterification conditions. See,

25 for instance, U.S. Patent Nos. 4,384,115 and 4,423,205 which are incorporated herein by reference in their entirety. Additional suitable cyclic aliphatic

carbonates are disclosed in U.S. Patent No. 4,747,850 which is also incorporated herein by reference in its entirety.

5 The term "viscosifier" refers to a suitable viscosifier for solid electrolytes. Viscosifiers include conventional viscosifiers such as those known to one of ordinary skill in the art. Suitable viscosifiers include film forming agents well known in the art which include, by way of example, polyethylene oxide, polypropylene oxide, copolymers thereof, and the like, having a number average molecular weight of at least about 100,000, polyvinylpyrrolidone, carboxymethylcellulose, and the like.

10 The term "toughening agent" refers to a solid particulate, compatible in solid electrolytes, that enhances the solid electrolyte's structural integrity and adherence to the anode and cathode surfaces. Suitable solid particulates include alumina (Al_2O_3), silica (SiO_2) and zeolites. Aluminum oxide is most preferred since it can also function as a basic sink material that removes or
15 neutralizes acids in the electrolyte. The basic sink material can be introduced in either the neutral or basic form because the surface of Al_2O_3 can be altered. Most likely, it is believed that the surface has the structure: $\text{Al}(\text{O})(\text{OH})$ with the number of OH groups determining its basicity. In the neutral or basic form, the OH group on the alumina will strongly attract the F of HF via a hydrogen
20 bond: $\text{Al}(\text{O})\text{OH} \cdots \text{FH}$. Thus, HF can be viewed as a sink because the HF reacts with or is "bonded" to the alumina and will not be released. The hydrogen bond between H and F is one of the strongest. The alumina effectively removes the HF from the system.

25 Zeolites suitable for use in the present invention include natural zeolites, such as hydrated silicates of aluminum and at least one of sodium and calcium, and synthetic zeolites, such as those made, for example, by a gel process (sodium silicate and alumina) and a clay process (kaolin), which form a matrix

to which the zeolite is added and involve the substitution of various rare-earth oxides. Suitable toughening agents also include non-reactive metal oxide materials, e.g., calcium oxide and magnesium oxide. Mixtures of different toughening agents can be used. The toughening agent is preferably in powder
5 form. Preferably, the size of the toughening agent ranges from about 0.1μ to about 10μ , more preferably from about 0.3μ to about 5μ , and most preferably from about 1μ to about 3μ .

Preferably the toughening agent comprises about 0.1 % (wt) to about 13 % (wt), more preferably about 2 % (wt) to about 8 % (wt), and most
10 preferably about 3 % (wt) to about 6 % (wt) of the electrolyte layer.

The term "electrolytic cell" or "electrochemical cell" refers to a composite containing an anode, a cathode and an ion-conducting electrolyte interposed therebetween.

The term "battery" refers to two or more electrochemical cells
15 electrically interconnected in an appropriate series/parallel arrangement to provide the required operating voltage and current levels.

The anode is typically comprised of a compatible anodic material which is any material which functions as an anode in a solid electrolytic cell. Such compatible anodic materials are well known in the art and include, by way of
20 example, lithium, lithium alloys, such as alloys of lithium with aluminum, mercury, manganese, iron, zinc, intercalation based anodes such as those employing carbon, tungsten oxides, and the like. Particularly preferred anodes include lithium intercalation anodes employing carbon materials such as graphite, cokes, mesocarbons, and the like. Such carbon intercalation based
25 anodes typically include a polymeric binder suitable for forming a bound porous composite having a molecular weight of from about 1,000 to 5,000,000.

Examples of suitable polymeric binders include EPDM (ethylene propylene diene termonomer), PVDF (polyvinylidene difluoride), EAA (ethylene acrylic acid copolymer), EVA (ethylene vinyl acetate copolymer), EAA/EVA copolymers, and the like. The anode also may include an electron conducting material such as carbon black.

The cathode is typically comprised of a compatible cathodic material (i.e., insertion compounds) which is any material which functions as a positive pole in a solid electrolytic cell. Such compatible cathodic materials are well known in the art and include, by way of example, transition metal oxides, sulfides, and selenides, including lithiated compounds thereof. Representative materials include cobalt oxides, manganese oxides, molybdenum oxides, vanadium oxides, sulfides of titanium, molybdenum and niobium, the various chromium oxides, copper oxides, lithiated cobalt oxides, e.g., LiCoO_2 , lithiated manganese oxides, e.g., LiMn_2O_4 , lithiated nickel oxides, e.g., LiNiO_2 , and mixtures thereof. Cathode-active material blends of $\text{Li}_x\text{Mn}_2\text{O}_4$ (spinel) is described in U.S. Patent 5,429,890 which is incorporated herein. The blends can also include $\text{Li}_x\text{Mn}_2\text{O}_4$ (spinel) and at least one lithiated metal oxide selected from Li_xNiO_2 and Li_xCoO_2 wherein $0 < x \leq 2$.

A preferred method of fabricating an electrochemical cell is described herein. In addition, methods for preparing solid electrochemical cells and batteries are described in the art, for example, in U.S. Patent 5,300,373, 5,316,556, 5,346,385, 5,262,253, 5,028,500, 4,472,487, and 4,668,595, which are all incorporated herein.

In one preferred embodiment, the cathode is prepared from a cathode paste which comprises from about 35 to 65 weight percent of a compatible cathodic material; from about 1 to 20 weight percent of an electroconductive agent; from about 0 to 20 weight percent of polyethylene oxide having a

number average molecular weight of at least 100,000. When the cathode includes V_6O_{13} , the cathode paste preferably further comprises from about 10 to 50 weight percent of solvent comprising a 10:1 to 1:4 (w/w) mixture of an organic carbonate and a glyme. The paste further comprises from about 5
5 weight percent to about 25 weight percent of the solid matrix forming monomer or partial polymer thereof. Also included is an ion conducting amount of an inorganic ion salt. Generally, the amount of the salt is from about 1 to about 25 weight percent. (All weight percents are based on the total weight of the cathode.)

10 The electrolyte composition typically comprises from about 5 to about 25 weight percent of the inorganic ion salt based on the total weight of the electrolyte; preferably, from about 10 to 20 weight percent; and even more preferably from about 10 to about 15 weight percent. The percentage of salt depends on the type of salt and electrolytic solvent employed.

15 The electrolyte composition typically comprises from 40 to about 80 weight percent electrolyte solvent (e.g., organic carbonate/glyme mixture) based on the total weight of the electrolyte; preferably from about 60 to about 80 weight percent; and even more preferably about 70 weight percent. The
20 electrolyte composition typically comprises from about 5 to about 30 weight percent of the solid polymeric matrix based on the total weight of the electrolyte; preferably from about 10 to about 20 weight percent. The electrolyte composition comprises from about 1 to about 15 weight percent toughening agent, based on the total weight of the electrolyte; preferably from about 5 to about 10 weight percent.

25 In a preferred embodiment, the electrolyte composition further comprises a small amount of a film forming agent. Suitable film forming agents are well known in the art and include, by way of example, polyethylene

oxide, polypropylene oxide, copolymers thereof, and the like, having a numbered average molecular weight of at least about 100,000. Preferably, the film forming agent is employed in an amount of about 1 to about 10 weight percent and more preferably at about 2.5 weight percent based on the total weight of the electrolyte composition.

EXAMPLE 1

A. The Cathode

The cathode may be prepared from a cathodic paste which, in turn, may be prepared from a cathode powder as follows:

10 i. Cathode Powder

The cathode powder can be prepared by combining 90.44 weight percent V_6O_{13} (prepared by heating ammonium metavanadate ($NH_4^+VO_3^-$) at 450°C for 16 hours under N_2 flow) and 9.56 weight percent of carbon (from Chevron Chemical Company, San Ramon, CA under the trade name of Shawinigan Black®). About 100 grams of the resulting mixture can be placed into a grinding machine (Attritor Model S-1 purchased from Union Process, Akron, Ohio) and ground for 45 minutes. Afterwards, the resulting mixture can be dried at about 260°C for 16 hours under vacuum which should provide a cathode powder having about 84.45 weight percent V_6O_{13} .

20 The above mixing procedure can be repeated to provide for a total of 292 grams of cathode powder.

ii. Cathode Paste

A cathode paste may be prepared by combining sufficient cathode powder which should provide for a final product having 45 weight percent V_6O_{13} .

Specifically, about 26.2 grams of unground carbon (from Chevron Chemical Company, San Ramon, CA under the trade name of Shawinigan Black®) can be combined in a glove box (under dry (< 10 ppm H_2O) argon at ambient temperature and pressure) with about 169.9 grams of a 4:1 w/w mixture of propylene carbonate/triglyme and the resulting composite mixed under dry argon and at ambient temperature and pressure on a double planetary mixer (Ross #2 mixer available from Charles Ross & Sons, Company, Hauppag, New York) at about 25 rpms until a paste is formed.

About 225.0 grams of a cathode powder prepared in a manner similar to that described above can be added to the mixer along with an additional 169.9 grams of the 4:1 mixture of propylene carbonate/triglyme and the resulting composite can be mixed under dry argon and at ambient temperature and pressure on a double planetary mixer at about 25 rpms until a dry paste is formed.

About 5 grams of polyethylene oxide (number average molecular weight about 600,000 available as Polyox WSR-205 from Union Carbide Chemicals and Plastics, Danbury, CT), about 42.5 grams of polyethylene glycol diacrylate (molecular weight about 500 available as SR-344 from Sartomer Company, Inc., Exton, PA) and containing less than about 50 ppm of inhibitor, and about 7.5 grams of ethoxylated trimethylpropane triacrylate (TMPEOTA) (molecular weight about 425 available as SR-454 from Sartomer Company, Inc., Exton, PA) and containing less than about 50 ppm of inhibitor can be added to about 169.9 grams of a 4:1 mixture of propylene carbonate/triglyme and this mixture added to the mixer.

The resulting slurry in the mixer can be heated at about $65^{\circ}C$ while mixing for 2 hours at 60 rpms to provide for the cathodic paste which would have the following approximate weight percent of components:

	V ₆ O ₁₃	45.00	weight percent
	Carbon	10.00	weight percent
	propylene carbonate	27.18	weight percent
	Triglyme	6.80	weight percent
5	Polyethylene glycol diacrylate	8.51	weight percent
	Ethoxylated trimethylpropane triacrylate ¹	1.51	weight percent
	Polyethylene oxide	1.00	weight percent

10 In an alternative embodiment, the requisite amounts of all of the cathodic materials other than the cathode powder can be combined to form a first mixture and this first mixture is combined with the cathode powder to form a second mixture. This second mixture can be then thoroughly mixed to provide for the cathode paste.

15 The cathode paste which can be prepared as above can be placed onto a sheet (about 1 mil (N-25 μ m) thick by 10 cm wide) of a roughened nickel on nickel current collector (available as CF18/NiT from Fukuda Metal Foil & Powder Company, Ltd., Kyoto, Japan). A Mylar cover sheet can then be placed over the paste and the paste spread to a thickness of about 75 microns
20 (μ m) with a conventional plate and roller system and cured by continuously passing the sheet through an electron beam apparatus (Electrocurtain, Energy Science Inc., Woburn, MA) at a voltage of about 175 kV and a current of about 12 mA and at a conveyor belt speed setting of 50 which provides a conveyor speed of about 3 in/sec. After curing, the Mylar sheet can be
25 removed which should provide for a solid cathode laminated to a nickel on nickel current collector.

30 ¹ Inhibitor may be removed from both the polyethylene glycol diacrylate and ethoxylated trimethylpropane triacrylate by contacting each of these compounds with an Inhibitor Remover available as Product No. 31,133-2 from Aldrich Chemical, Milwaukee, Wisconsin, which results in less than 50 ppm of inhibitor in the product.

B. Electrolyte

The electrolyte may be prepared by first combining 52 grams of propylene carbonate, 13 grams of triglyme and 17 grams of urethane acrylate (available as Photomer 6140 from Henkel Corporation, Coating and Chemicals Division, Ambler, PA). The propylene carbonate/triglyme/urethane acrylate mixture can be dried over molecular sieves (Grade 514, 4Å, 8-12 mesh, available from W. R. Grace, Baltimore, MD) to remove water.

This solution can then be combined with 3 grams of polyethylene oxide (weight average molecular weight about 600,000 available as Polyox WSR-205 from Union Carbide Chemicals and Plastics, Danbury, CT). The mixture can then be thoroughly mixed with the same laboratory mixer at heating until a temperature of about 65°C is reached and then cooled to ambient temperature over at least a 2 hour period while stirring is maintained.

Once the polyethylene oxide is dispersed and dissolved, 5 grams of LiAsF₆ (available from FMC Corporation Lithium Division, Bessemer City, NC, as Lectrosalt®), 5 grams of Li(OCH₂CH₂)₃OCH₃ and 5 grams of alumina can be added while stirring with a laboratory mixer (Yamato Model LR41B, available from Fisher Scientific, Santa Clara, CA). The Li(OCH₂CH₂)₃OCH₃ salt can be prepared from a 1:1 molar ratio n-butyl lithium and the monomethylether of triethylene glycol in tetrahydrofuran at -78°C. Other aprotic solvents could also be used.

The resulting 100 gram mixture would contain the following weight percent of components:

25	Propylene carbonate	52 weight percent
	Triglyme	13 weight percent
	Urethane acrylate (Photomer 6140)	17 weight percent
	LiAsF ₆	5 weight percent
	Li(OCH ₂ CH ₂) ₃ OCH ₃	5 weight percent

Polyethylene oxide
Alumina

3 weight percent
5 weight percent.

Afterwards, the electrolyte mixture can then be coated by a conventional knife blade to a thickness of about 50 μm onto the surface of the cathode sheet prepared as above (on the side opposite that of the current collector) but without the Mylar covering. The electrolyte can then be cured by continuously passing the sheet through an electron beam apparatus (Electrocurtain, Energy Science Inc., Woburn, MA) at a voltage of about 175 kV and a current of about 1.0 mA and at a conveyor speed setting of 50 which provides for a conveyor speed of about 1 cm/sec. After curing, a composite is recovered which should contain a solid electrolyte laminated to a solid cathode which, in turn, is laminated to a nickel on nickel current collector.

C. Anode

The anode may comprise a sheet of lithium foil (about 76 μm thick) which is commercially available from FMC Corporation Lithium Division, Bessemer City, NC.

D. The Solid Battery

A solid battery may be prepared by first preparing a cathodic paste as described above which can be spread onto a substrate, e.g., a current collector, and then cured to provide the cathode. An electrolyte composition as described above can then be placed onto the cathode surface and cured to provide for the solid electrolyte. Then, the anode can be laminated onto the solid electrolyte to provide for the solid battery.

EXAMPLE 2

A solid electrolytic cell can be prepared by first preparing a cathodic paste which is spread onto a current collector and cured to provide for the

cathode. An electrolyte solution can be placed onto the cathode surface and cured to provide for the solid electrolyte composition. Then, the anode can be laminated onto the solid electrolyte composition to provide for a solid electrolytic cell. The specifics of this construction are as follows:

5 A. The Current Collector

The current collector is a sheet of aluminum foil having a layer of adhesion promoter attached to the surface of the foil which should contact the cathode so as to form a composite having a sheet of aluminum foil, a cathode and a layer of adhesion promoter interposed therebetween.

10 Specifically, the adhesion promoter layer can be prepared as a dispersed colloidal solution by one of two methods. The first preparation of this colloidal solution for this example can be as follows:

15 84.4 weight percent of carbon powder
 (Shawinigan Black™-- available from Chevron Chemical
 Company, San Ramon, CA)

20 337.6 weight percent of a 25 weight percent
 solution of polyacrylic acid (a reported average molecular weight
 of about 90,000, commercially available from Aldrich Chemical
 Company -- contains about 84.4 grams polyacrylic acid and
 253.2 grams water)

 578.0 weight percent of isopropanol

25 The carbon powder and isopropanol can be combined with mixing in a conventional high shear colloid mill mixer (Ebenbach-type colloid mill) until the carbon is uniformly dispersed and the carbon particle size is smaller than 10 microns. At this point, the 25 weight percent solution of polyacrylic acid can be added to the solution and mixed for approximately 15 minutes. The resulting mixture can be placed on the coating head and roll coated with a Meyer rod onto a sheet of aluminum foil (9 inches wide and about 0.0005

inches thick). After application, the solution/foil can be contacted with a Mylar wipe (about 0.002 inches thick by about 2 inches and by about 9 inches wide -- the entire width of aluminum foil). The wipe can be flexibly engaged with the foil, i.e., the wipe contacts the foil, to redistribute the solution so as to provide for a substantially uniform coating. Evaporation of the solvents, i.e., water and isopropanol, via a conventional gas-fired oven can provide for an electrically conducting adhesion-promoter layer of about 6 microns in thickness or about 3×10^{-4} grams per cm^2 . The aluminum foil can be cut to about 8 inches wide by removing approximately $\frac{1}{2}$ inch from either side by the use of a conventional slit-ter so as to remove any uneven edges.

In order to further remove the aprotic solvent from this layer, the foil can be redried. In particular, the foil can be wound up and a copper support can be placed through the roll's cavity. The roll can be hung overnight from the support in a vacuum oven maintained at about 130°C . Afterwards, the roll can be removed. In order to avoid absorption of moisture from the atmosphere, the roll can be preferably stored in a desiccator or other similar anhydrous environment to minimize atmospheric moisture content until the cathode paste is ready for application onto this roll.

The second preparation of this colloidal solution can comprise mixing 25 lbs of carbon powder (Shawinigan BlackTM -- available from Chevron Chemical Company, San Ramon, CA) with 100 lbs of a 25 weight percent solution of polyacrylic acid (average molecular weight of about 240,000, commercially available from BF Goodrich, Cleveland, Ohio, as Good-Rite K702 -- contains about 25 lbs polyacrylic acid and 75 lbs water) and with 18.5 lbs of isopropanol. Stirring can be done in a 30 gallon polyethylene drum with a gear-motor mixer, e.g., Lighting Labmaster Mixer, model XJ-43, available from Cole-Parmer Instruments Co., Niles, Illinois, at 720 rpm with two 5 inch diameter A310-type propellers mounted on a single shaft. This procedure can

wet down the carbon and eliminate any further dust problem. The resulting weight of the mixture should be about 143.5 lbs and contain some "lumps".

5 The mixture can be further mixed with an ink mill which consists of three steel rollers almost in contact with each other, turning at 275, 300, and 325 rpms, respectively. This high shear operation allows particles that are sufficiently small to pass directly through the rollers. Those that do not pass through the rollers can continue to mix in the ink mill until they are small enough to pass through these rollers. When the mixing is complete, the carbon powder is completely dispersed. A Hegman fineness of grind gauge (available from Paul N. Gardner Co., Pompano Beach, FL) should indicate that the particles are 4-6 μm with the occasional 12.5 μm particles. The mixture can be stored for well over 1 month without the carbon settling out or reagglomerating.

15 When this composition is to be used to coat the current collector, an additional 55.5 lbs of isopropanol can be mixed into the composition working with 5 gallon batches in a plastic pail using an air powered shaft mixer (Dayton model 42231 available from Granger Supply Co., San Jose, CA) with a 4 inch diameter Jiffy-Mixer brand impeller (such as an impeller available as Catalog No. G-04541-20 from Cole Parmer Instrument Co., Niles, Illinois). The composition can be gear pumped through a 25 μm cloth filter, e.g., So-Clean Filter Systems, American Felt and Filter Company, Newburgh, NY, and Meyer-rod coated as described above.

B. The Cathode

25 The cathode can be prepared from a cathodic paste which, in turn, can be prepared from a cathode powder as follows:

i. Cathode Powder

The cathode powder can be prepared by combining 90.44 weight percent V_6O_{13} [prepared by heating ammonium metavanadate ($NH_4^+VO_3^-$) at $450^\circ C$ for 16 hours under N_2 flow] and 9.56 weight percent of carbon (from Chevron Chemical Company, San Ramon, CA under the tradename of Shawinigan BlackTM). About 100 grams of the resulting mixture can be placed into a grinding machine (Attritor Model S-1 purchased from Union Process, Akron, Ohio) and ground for 30 minutes. Afterwards, the resulting mixture can be dried at about $260^\circ C$ for 21 hours.

ii. Cathode Paste

A cathode paste can be prepared by combining sufficient cathode powder which should provide for a final product having 45 weight percent V_6O_{13} .

Specifically, 171.6 grams of a 4:1 weight ratio of propylene carbonate:triglyme can be combined with 42.9 grams of polyethylene glycol diacrylate (molecular weight about 400 available as SR-344 from Sartomer Company, Inc., Exton, PA), and about 7.6 grams of ethoxylated trimethylolpropane triacrylate (TMPEOTA) (molecular weight about 450 available as SR-454 from Sartomer Company, Inc., Exton, PA) in a double planetary mixer (Ross #2 mixer available from Charles Ross & Sons, Company, Hauppauge, New York).

A propeller mixture can be inserted into the double planetary mixer and the resulting mixture can be stirred at 150 rpms until homogeneous. The resulting solution can be passed through sodiated 4\AA molecular sieves. The solution can be returned to a double planetary mixer equipped with the propeller mixer and about 5 grams of polyethylene oxide (number average molecular weight about 600,000 available as Polyox WSR-205 from Union Carbide Chemicals and Plastics, Danbury, CT) can be added to the solution vortex by a mini-sieve such as a 25 mesh mini-sieve commercially available as

Order No. 57333-965 from VWR Scientific, San Francisco, CA.

The solution can be heated while stirring until the temperature of the solution reaches 65°C. At this point, stirring can be continued until the solution is completely clear. The propeller blade can be removed and the cathode powder prepared as above can be added as well as an additional 28.71 grams of unground carbon (from Chevron Chemical Company, San Ramon, CA under the tradename of Shawinigan Black™). The resulting mixture can be mixed at a rate of 7.5 cycles per second for 30 minutes in the double planetary mixer. During this mixing the temperature can slowly increase to a maximum of 73°C. At this point, the mixing can be reduced to 1 cycle per second and the mixture can be slowly cooled to a temperature of from 40°C to 48°C, e.g., about 45°C. The resulting cathode paste can be maintained at this temperature until just prior to application onto the current collector.

The resulting cathode paste can have the following approximate weight percent of components:

15	V ₆ O ₁₃	45 weight percent
	Carbon	10 weight percent
	4:1 propylene carbonate/tri-glyme	34 weight percent
	polyethylene oxide	1 weight percent
20	polyethylene glycol diacrylate	8.5 weight percent
	ethoxylated trimethylol-propane triacrylate	1.5 weight percent

In an alternative embodiment, the requisite amounts of all of the solid components can be added directly to the combined liquid components. In this regard, mixing speeds can be adjusted to account for the amount of the material mixed and size of vessel used to prepare the cathode paste. Such adjustments are well known to the skilled artisan.

In order to enhance the coatability of the carbon paste onto the current

collector, it may be desirable to heat the paste to a temperature of from about 60°C to about 130°C and more preferably, from about 80°C to about 90°C and for a period of time of from about 0.1 to about 2 hours, more preferably, from about 0.1 to 1 hour and even more preferably from about 0.2 to 1 hour. A
5 particularly preferred combination is to heat the paste at from about 80°C to about 90°C for about 0.33 to about 0.5 hours.

During this heating step, there is no need to stir or mix the paste although such stirring or mixing may be conducted during this step. However, the only requirement is that the composition be heated during this period. In
10 this regard, the composition to be heated should have a volume to surface area ratio such that the entire mass is heated during the heating step.

The cathode paste noted above can be placed onto the adhesion layer of the current collector described above by extrusion at a temperature of from about 45° to about 48°C. A Mylar cover sheet can be placed over the paste
15 and the paste can be spread to a thickness of about 90 microns (μm) with a conventional plate and roller system and can be cured by continuously passing the sheet through an electron beam apparatus (Electro-curtain, Energy Science Inc., Woburn, MA) at a voltage of about 175 kV and a current of about 1.0 mA and at a rate of about 1 cm/sec. After curing, the Mylar sheet can be
20 removed to provide for a solid cathode laminated to the aluminum current collector described above.

C. Electrolyte

48 grams of propylene carbonate, 12 grams of triglyme, and 17 grams of urethane acrylate (Photomer 6140, available from Henkel Corp., Coating and
25 Chemical Division, Ambler, PA) can be combined at room temperature until homogeneous. The resulting solution can be passed through a column of 4Å sodiated molecular sieves to remove water and then can be mixed at room

temperature until homogeneous.

At this point, 4 grams of polyethylene oxide having a number average molecular weight of about 600,000 (available as Polyox WSR-205 from Union Carbide Chemicals and Plastics, Danbury, CT) and 10 grams of silica can be added to the solution and then dispersed while stirring with a magnetic stirrer over a period of about 120 minutes. After dispersion, the solution can be heated to between 60°C and 65°C with stirring until the viscosifier dissolves. The solution can be cooled to a temperature of between 45° and 48°C, a thermocouple can be placed at the edge of the vortex created by the magnetic stirrer to monitor solution temperature, and then 9 grams of LiPF₆ can be added to the solution over a 120 minute period while thoroughly mixing to ensure a substantially uniform temperature profile throughout the solution. Cooling can be applied as necessary to maintain the temperature of the solution between 45° and 48°C.

In one embodiment, the polyethylene oxide can be added to the solution via a mini-sieve such as a 25 mesh mini-sieve commercially available as Order No. 57333-965 from VWR Scientific, San Francisco, CA.

The resulting solution should contain the following:

	<u>Component</u>	<u>Amount</u>
20	Propylene Carbonate	48 g
	Triglyme	12 g
	Urethane Acrylate	17 g
	LiPF ₆	9 g
	PEO	4 g
25	Silica	10 g
	Total	100 g

This solution can be degassed to provide for an electrolyte solution wherein little, if any, of the LiPF_6 salt decomposes.

Optionally, solutions which can be produced as above and which contain the prepolymer, the polyethylene oxide, the toughening agent, the electrolyte solvent and the LiPF_6 salt can be filtered to remove any solid particles or gels remaining in the solution. One suitable filter device is a sintered stainless steel screen having a pore size between 1 and 50 μm at 100% efficiency. Smaller screens may be required depending on the size of the toughening agent particles.

Alternatively, the electrolyte solution can be prepared in the following manner. Specifically, in this example, the mixing procedure can be conducted using the following weight percent of components:

	Propylene Carbonate	44 weight percent
	Triglyme	11 weight percent
15	Urethane Acrylate ^b	20 weight percent
	LiPF_6	10 weight percent
	PEO ^c	5 weight percent
	Silica	10 weight percent

^b (Photomer 6140, available from Henkel Corp., Coating and Chemical Division, Ambler, PA)

^c polyethylene oxide having a number average molecular weight of about 600,000 (available as Polyox WSR-205 from Union Carbide Chemicals and Plastics, Danbury, CT)

The mixing procedure employs the following steps:

1. Check the moisture level of the urethane acrylate. If the moisture level is less than 100 ppm water, proceed to step 2. If not, then first dissolve the urethane acrylate at room temperature, $<30^\circ\text{C}$, in the propylene

carbonate and triglyme and dry the solution over sodiated 4Å molecular sieves (Grade 514, 8-12 Mesh from Schoofs Inc., Moraga, CA) and then proceed to step 4.

2. Dry the propylene carbonate and triglyme over sodiated 4Å molecular sieves (Grade 514, 8-12 Mesh from Schoofs Inc., Moraga, CA).

3. At room temperature, <30°C, add the urethane acrylate to the solvent prepared in step 2. Stir at 300 rpm until the resin is completely dissolved. The solution should be clear and colorless.

4. Dry and then sift the polyethylene oxide through a 25 mesh mini-sieve commercially available as Order No. 57333-965 from VWR Scientific, San Francisco, CA. While stirring at 300 rpm, add the dried and pre-sifted polyethylene oxide slowing to the solution. The polyethylene oxide should be sifted into the center of the vortex formed by the stirring means over a 30 minute period. Addition of the polyethylene oxide should be dispersive and, during addition, the temperature should be maintained at room temperature (<30°C).

5. After final addition of the polyethylene oxide, add the silica and stir an additional 30 minutes to ensure that the viscosifier is substantially dispersed.

6. Heat the mixture to 68°C to 75°C and stir until the viscosifier has melted and the solution has become transparent to light yellow in color. Optionally, in this step, the mixture is heated to 65°C to 68°C.

7. Cool the solution produced in step 6 and when the temperature of the solution reaches 40°C, add the LiPF₆ salt very slowly making sure that the maximum temperature does not exceed 55°C.

8. After the final addition of the LiPF₆ salt, stir for an additional 30 minutes, degas, and let sit overnight and cool.

9. Filter the solution through a sintered stainless steel screen having a pore size between 1 and 50 μm at 100% efficiency.

At all times, the temperature of the solution should be monitored with a thermocouple which should be placed in the vortex formed by the mixer.

Afterwards, the electrolyte mixture is then coated by a conventional knife blade to a thickness of about 50 μm onto the surface of the cathode sheet prepared as above (on the side opposite that of the current collector) but without the Mylar covering. The electrolyte should be cured by continuously passing the sheet through an electron beam apparatus (Electrocurtain, Energy Science Inc., Woburn, MA) at a voltage of about 175 kV and a current of about 1.0 mA and at a conveyor speed setting of 50 which provides for a conveyor speed of about 1 cm/sec. After curing, a composite can be recovered which contains a solid electrolyte laminated to a solid cathode.

D. Anode

The anode can comprise a sheet of lithium foil (about 76 μm thick) which is commercially available from FMC Corporation Lithium Division, Bessemer City, North Carolina.

E. The Solid Electrolytic Cell

A sheet comprising a solid battery can be prepared by laminating the lithium foil anode to the surface of the electrolyte in the sheet produced in step C above. Lamination can be accomplished by minimal pressure.

20

EXAMPLE 3

Experiments were conducted which demonstrated that adding alumina to electrolytes did not adversely effect its viscosity or ionic conductivity. Alumina was added to two electrolyte solutions in accordance with the procedures set forth in Example 2. The two electrolyte solutions are designated "premix" and

"standard electrolyte" herein and the components for each solution are set forth in the following table. The alumina used was in powder form with an average size of 1.5 micron. (The alumina was from Fujimi Corp., Elmhurst, IL.)

Premix		Standard Electrolyte	
	Weight %		Weight %
5			
Propylene Carbonate	58.00	Propylene Carbonate	56.51
Triglyme	14.50	Triglyme	14.13
Urethane Acrylate	18.02	Urethane Acrylate	17.56
LiPF ₆	9.48	LiPF ₆	9.24
		PEO	2.56
10			
Total	100%	Total	100%

As is apparent, the components of the premix and the standard electrolyte are the same except that the latter contains polyethylene oxide which is a viscosifier. Alumina was added to the formulations to produce 5% and 10% (wt) alumina mixtures. Viscosity measurements indicate that the addition of alumina had negligible effect on the viscosity.

The conductivities of the uncured electrolyte mixtures containing alumina were measured and the results are as follows:

	σ/mScm^{-1}
20	
Premix 5%	3.19
Premix 10%	3.09
Std. Electrolyte 5%	2.82
Std. Electrolyte 10%	2.86
Std. Electrolyte (without toughening agent)	3.00

25 The data indicate that the electrolytes maintained good conductivity with the addition of alumina.

The following illustrates a method of fabricating an electrolytic cell wherein the solid polymeric matrix is formed by a casting process. Examples 4A and 4B describe the process of preparing the anode slurry and cathode slurry, respectively. Example 4C describes the procedures for assembling the electrolytic cell.

EXAMPLE 4A

The anode current collector employed was a sheet of expanded copper metal that is about 50 μ m thick. It is available under the designation 2Cu5-125 (flatten) from Delker Corp., Branford, Connecticut.

The anode slurry was prepared as follows:

A polymer mixture comprising a copolymer of polyvinylidene difluoride (PVDF) and hexafluoropropylene (HFP) was prepared by mixing 6.8 grams of the copolymer in 20 grams of acetone. The copolymer (ave. MW 125K) was Kynar Flex 2801™ from Elf Atochem North America, in Philadelphia, Pennsylvania. The mixture was stirred for about 24 hours in a milling jar available from VWR Scientific, in San Francisco, California, model H-04172-00. The copolymer functions as a binder for the carbon in the anode.

A graphite mixture was prepared separately by first adding 23.4 grams of graphite into 0.9 grams of carbon black into a solution containing 60 grams acetone, and 10.5 grams dibutyl phthalate. A preferred graphite comprises a 50:50 (by weight) mixture of a synthetic graphite available under the designation SFG-15™ from Lonza G & T, Ltd. (Sins, Switzerland) and graphitized mesocarbon microbeads available under the designation MCMB25-28™ from Osaka Gas Co., Japan. A preferred carbon black is available under the designation Super P™ from M.M.M. Carbon, Willebroek, Belgium. The

graphite mixture was then vigorously mixed in a high shear mixer until a substantially homogeneous blend was formed. A suitable mixer is available from Ross Model ME100DLX, Hauppauge, New York, operating at its highest setting (about 10,000 RPM) for 30 minutes.

5 The anode slurry was prepared by mixing the polymer mixture and the graphite mixture together under low shear conditions to form the anode slurry wherein the components are well mixed. A portion of the acetone is allowed to evaporate from the slurry before it was laminated onto each side of the current collector. Anode films form when the remaining portion of the acetone
10 evaporates.

EXAMPLE 4B

The cathode current collector employed was a sheet of expanded aluminum that is about 50 μm thick. The aluminum grid is available under the designation 2A1 077 from Delkor Corp.

15 The cathode slurry was prepared as follows:

A polymer mixture comprising a copolymer of polyvinylidenedifluoride (PVDF) and hexafluoropropylene (HFP) was prepared by mixing 4.4 grams of the copolymer in 15 ml of acetone. The copolymer was Kynar Flex 2801™. The mixture was stirred for about 24 hours in a milling jar.
20

A cathode mixture was prepared separately by first adding 28.9 grams of a cathode-active material comprising $\text{Li}_x\text{Mn}_2\text{O}_4$ (spinel) ($0 \leq x < 2$), 2.4 grams of carbon black (Super P™) into a solution containing 60 grams acetone, and 8.7 grams dibutyl phthalate. The mixture was then vigorously mixed in a high
25 shear mixer until a substantially homogeneous blend was formed.

The cathode slurry was prepared by mixing the polymer mixture and the

cathode mixture together under low shear conditions to form the cathode slurry wherein the components are well mixed. A portion of the acetone is allowed to evaporate from the slurry before it was laminated onto each side of the current collector. Cathode films form when the remaining portion of the acetone
5 evaporates.

The above anode and cathode films were formed directly on the current collector by laminating the slurry mixtures onto the current collector surfaces. Alternatively, each film can be prepared by first casting a slurry onto a substrate or carrier web and allowing the solvent to evaporate thus leaving the
10 film. Thereafter, the films can be laminated onto each side of the current collector.

EXAMPLE 4C

A solid electrochemical cell precursor is prepared by positioning a polymeric matrix between the anode and cathode and thereafter laminating the
15 structures under moderate pressure and/or temperature (e.g., 130°C). The pressure and temperature employed will depend on the polymer(s) forming the matrix. The polymeric matrix is formed by casting a slurry comprising acetone, dibutyl phthalate, silanized fumed SiO₂, and a PVDF/HFP copolymer onto a carrier web or other substrate and allowing the acetone to evaporate. No
20 curing by radiation is required. The SiO₂ is a filler which imparts toughness and strength to the film. In addition, it is believed that the SiO₂ assists the activation process by creating physico-chemical conditions such that the electrolyte solution quickly and completely fills the pores created by the extraction of the dibutyl phthalate. Preferably, the slurry is mixed under low
25 shear conditions as not to degrade the copolymer.

Next the dibutyl phthalate, a plasticizer, is extracted from the precursor or plurality of electrochemical cell precursors in the case of a battery. Extraction

can be accomplished using conventional organic liquid solvents such as diethyl ether or by a supercritical fluid which includes, for example, a gas compressed and heated to either supercritical or subcritical conditions to achieve liquid-like densities. Supercritical fluids are known in the art. See, for example, U.S. Patents 5,013,366, 5,267,455, 4,219,313, 4,012,194, and 3,969,196, which are incorporated herein. A preferred supercritical fluid is carbon dioxide.

Following extraction, the precursor is then packaged in moisture-impermeable material which is described, for example, in U.S. Patent 5,326,653 which is incorporated herein. The precursor is then activated by the addition of an inorganic salt and electrolyte solvent. Extraction and activation preferably take place in an inert (e.g., argon) atmosphere. Finally, the packaging is sealed. In assembling the components, the anode precursor, cathode precursor, and polymeric electrolyte precursor are preferably fused together by the application of moderate pressure and heat (e.g., 130° C) to cause the polymeric components of the anode and cathode precursors to adhere to the polymeric layer.

Although only preferred embodiments of the invention are specifically disclosed and described above, it will be appreciated that many modifications and variations of the present invention are possible in light of the above teachings and within the purview of the appended claims without departing from the spirit and intended scope of the invention.

WHAT IS CLAIMED IS:

1. An electrolytic cell which comprises:
an anode; a cathode; and interposed therebetween a solid, solvent-containing electrolyte which comprises:
 - 5 a solid polymeric matrix;
 - a salt;
 - a solvent; and
 - a toughening agent.
2. The electrolytic cell of Claim 1, wherein said toughening agent is
10 selected from the group consisting of alumina, silica, zeolite, calcium oxide, magnesium oxide, and mixtures thereof.
3. The electrolytic cell of Claim 1, wherein the toughening agent
comprises from about 1% to about 15% by weight of the solid electrolyte.
4. The electrolytic cell of Claim 2, wherein the toughening agent
15 comprises from about 3 % to about 8 % by weight of the solid electrolyte.
5. The electrolytic cell of Claim 1 wherein the anode comprises an
intercalation based anode comprising carbon.
6. The electrolytic cell of Claim 1 wherein the cathode comprises
material selected from the group consisting of vanadium oxide, lithiated
20 manganese oxide, lithiated nickel oxide, lithiated cobalt oxide, and mixtures thereof.
7. The electrolytic cell of Claim 6, wherein the anode comprises an
intercalation based anode comprising carbon.

8. The electrolytic cell of Claim 1 wherein said toughening agent comprises alumina.
9. The electrolytic cell of Claim 8, wherein the toughening agent comprises from about 3 % to about 8 % by weight of the solid electrolyte.
- 5 10. The electrolyte cell of Claim 8 wherein the salt is LiPF_6 .
11. The electrolytic cell of Claim 1 wherein said toughening agent comprises silica.
12. A process for preparing an electrolytic cell which comprises:
- 10 (a) providing a cathode;
- (b) providing an anode;
- (c) forming an electrolyte between the cathode and anode wherein the electrolyte comprises:
- 15 a polymeric matrix,
- a salt,
- a solvent; and
- a toughening agent.
13. The process of Claim 12, wherein said toughening agent is selected from the group consisting of alumina, silica, zeolite, calcium oxide, magnesium oxide, and mixtures thereof.
- 20 14. The process of Claim 12, wherein the toughening agent comprises from about 1% to about 15% by weight of the electrolyte.
15. The process of Claim 13, wherein the toughening agent comprises

from about 3% to about 8% by weight of the electrolyte.

16. The process of Claim 12 wherein the anode comprises an intercalation based anode comprising carbon.

5 17. The process of Claim 12 wherein the cathode comprises material selected from the group consisting of vanadium oxide, lithiated manganese oxide, lithiated nickel oxide, lithiated cobalt oxide, and mixtures thereof.

18. The process of Claim 17 wherein the anode comprises an intercalation based anode comprising carbon.

10 19. The process of Claim 12 wherein said toughening agent comprises alumina.

20. The process of Claim 19 wherein the toughening agent comprises from about 3% to about 8% by weight of the electrolyte.

21. The process of Claim 19 wherein the salt is LiPF_6 .

15 22. The process of Claim 12 wherein the toughening agent comprises SiO_2 .

20 23. A method for removing HF from the electrolyte layer of an electrochemical cell that includes an anode, a cathode, and interposed therebetween a solvent-containing electrolyte layer comprising LiPF_6 and a solvent, which comprises adding an effective amount of alumina to the electrolyte layer to react with HF that is dissolved in the solvent.

24. The method of Claim 23 wherein the alumina comprises about 3% to

about 8% by weight of the electrolyte layer.

25. The method of Claim 24 wherein electrolyte layer comprises a polymeric matrix.

26. The method of Claim 23 wherein the alumina forms hydrogen bonds
5 with the HF.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/18240

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :H01M 10/36

US CL :429/192

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 429/192

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
none

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS: solid (w) polymer, alumina, silica, zeolite #, electrolyte

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	Second International Symposium on Polymer Electrolytes, 1990, W. Wieczorek et al, "Mixed Phase said Electrolytes based on Peo Systems", see pages 339-346.	1-26
Y	US, 4,990,413 A (LEE ET AL) 05 February 1991 (05-02-91), see columns 5-6.	1-26
Y,P	US 5,616,437 A (GAO) 01 April 1997 (01-04-97), see columns. 1, 3, 4, 6, 8.	1-26



Further documents are listed in the continuation of Box C.



See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
E earlier document published on or after the international filing date	*Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*A* document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means	
P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

23 JANUARY 1998

Date of mailing of the international search report

26 FEB 1998

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231

Facsimile No. (703) 305-3230

Authorized officer

JOHN S. MAPLES

Telephone No. (703) 308-0661

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US97/18240

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of Item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☒ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

Form PCT/ISA/210 (continuation of first sheet(1))(July 1992)*

THIS PAGE BLANK (USPTO)